

QUALITY SYSTEM

EC-CERTIFICATE

Directive 93/42/EEC

Manufacturer: Suunto Oy
Tammiston kauppatie 7A
FI-01510 Vantaa
FINLAND

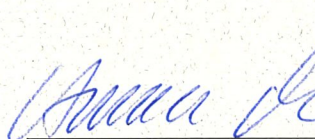
Coverage of Certificate: Design, manufacture and final inspection

Product category: Medical device accessories for measurement
of electrocardiogram and movement signals

Valid until: 27th May 2024

The manufacturer's quality system for the design, manufacture and final inspection of the aforesaid product category has been evaluated and meets the provisions of Council Directive 93/42/EEC as set out in Annex II Section 3. This approval is valid until the expiry date provided that the manufacturer fulfils the obligations imposed by Annex II in Directive 93/42/EEC. Products covered by the certificate are specified in the attachment(s).

Valid from: 4th December 2020





Anniina Mäkelä

Certificate no.
C-01-1205-754-20


Satu Rajala

Notified Body no. 0537:
Eurofins Expert Services
Kivimiehentie 4
FI-02150 ESPOO, FINLAND

Attachment 1 to the Certificate number: C-01-1205-754-20

Manufacturer:	Suunto Oy Tammiston kauppatie 7A, FI-01510 Vantaa Finland						
Activity and product category:	Design, manufacture and final inspection of medical device accessories for measurement of electrocardiogram and movement signals GMDN-code(s): 12391						
Products:	<p>The certificate covers the following products:</p> <table border="1"> <thead> <tr> <th><i>Name</i></th> <th><i>Brand name</i></th> <th><i>Class</i></th> </tr> </thead> <tbody> <tr> <td>Movesense MD</td> <td>Suunto</td> <td>IIa</td> </tr> </tbody> </table>	<i>Name</i>	<i>Brand name</i>	<i>Class</i>	Movesense MD	Suunto	IIa
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Movesense MD	Suunto	IIa					
Date:	Valid from: 4 th December 2020						
	 Anniina Mäkelä						
	  Satu Rajala						

Eurofins Expert Services is Notified Body no. 0537 under Council Directive 93/42/EEC.

EUROFINS EXPERT SERVICES

 Medical Devices
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Business ID FI22975132